

FILE 'USPATFULL' ENTERED AT 12:59:16 ON 20 AUG 2002  
CA INDEXING COPYRIGHT (C) 2002 AMERICAN CHEMICAL SOCIETY (ACS)

=> s l2 or hydroxyapatite or (hydroxy apatite) or l3 or (calcium phosphate)  
or acp

L5 81733 L2 OR HYDROXYAPATITE OR (HYDROXY APATITE) OR L3 OR (CALCIUM  
PHOSPHATE) OR ACP

=> s l2 or hydroxyapatite or (hydroxy apatite) or l3 or (calcium phosphate)  
or cap

<-----User Break----->

u

=> d 14

YOU HAVE REQUESTED DATA FROM FILE 'REGISTRY' - CONTINUE? (Y)/N:y

L4 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS

RN 33069-62-4 REGISTRY

CN Benzenepropanoic acid, .beta.- (benzoylamino) -.alpha.-hydroxy-,

(2aR,4S,4aS,6R,9S,11S,12S,12aR,12bS)-6,12b-bis(acetoxy)-12-(benzoyloxy)-  
2a,3,4,4a,5,6,9,10,11,12,12a,12b-dodecahydro-4,11-dihydroxy-4a,8,13,13-  
tetramethyl-5-oxo-7,11-methano-1H-cyclodeca[3,4]benz[1,2-b]oxet-9-yl  
ester, (.alpha.R,.beta.S)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 7,11-Methano-1H-cyclodeca[3,4]benz[1,2-b]oxete, benzenepropanoic acid  
deriv.

CN Benzenepropanoic acid, .beta.- (benzoylamino) -.alpha.-hydroxy-,  
6,12b-bis(acetoxy)-12-(benzoyloxy)-2a,3,4,4a,5,6,9,10,11,12,12a,12b-  
dodecahydro-4,11-dihydroxy-4a,8,13,13-tetramethyl-5-oxo-7,11-methano-1H-  
cyclodeca[3,4]benz[1,2-b]oxet-9-yl ester, [2aR-

[2a.alpha.,4.beta.,4a.beta.,6.beta.,9.alpha.(.alpha.R\*,.beta.S\*),11.alpha.  
,12.alpha.,12a.alpha.,12b.alpha.]-

CN Tax-11-en-9-one,

5.beta.,20-epoxy-1,2.alpha.,4,7.beta.,10.beta.,13.alpha.-  
hexahydroxy-, 4,10-diacetate 2-benzoate 13-ester with

(2R,3S)-N-benzoyl-3-  
phenylisoserine (8CI)

OTHER NAMES:

CN ABI 007

CN BMS 181339-01

CN NSC 125973

CN Paclitaxel

CN Plaxicel

CN Taxol

CN Taxol A

CN Yewtaxan

FS STEREOSEARCH

MF C47 H51 N O14

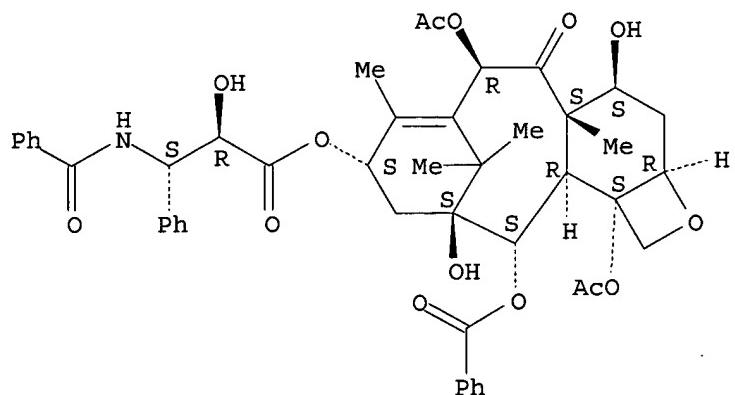
CI COM

LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN\*,  
BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB,  
CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, DDFU, DETHERM\*,  
DIOGENES, DRUGNL, DRUGPAT, DRUGU, DRUGUPDATES, EMBASE, HSDB\*, IFICDB,  
IFIUDB, IPA, MEDLINE, MRCK\*, MSDS-OHS, NAPRALERT, PHAR, PHARMASEARCH,  
PIRA, PROMT, RTECS\*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL,

VETU

(\*File contains numerically searchable property data)

Absolute stereochemistry. Rotation (-).



6294 REFERENCES IN FILE CA (1967 TO DATE)

338 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

6328 REFERENCES IN FILE CAPLUS (1967 TO DATE)

COPYRIGHT 2002 ACS  
AN 2002:408483 CAPLUS  
DN 137:10969  
TI Chemotherapeutic composition using nanocrystalline calcium phosphate  
paste  
IN Lee, Dosuk D.; Aiolova, Maria  
PA Etex Corporation, USA  
SO PCT Int. Appl., 50 pp.  
CODEN: PIXXD2  
DT Patent  
LA English  
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002041844	A2	20020530	WO 2001-US51419	20011019
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG		

PRAI US 2000-693120 A2 20001020

L9 ANSWER 2 OF 3 USPATFULL  
AN 2002:24062 USPATFULL  
TI Apatite-coated solid composition  
IN Saito, Kazuhiro, Suita, JAPAN  
Hoshino, Tetsuo, Osaka, JAPAN  
PA Takeda Chemical Industries, Ltd., Osaka, JAPAN (non-U.S. corporation)  
PI US 6344209 B1 20020205  
WO 9847485 19981029  
AI US 1999-403414 19991020 (9)  
WO 1998-JP1870 19980423  
19991020 PCT 371 date  
PRAI JP 1997-106918 19970424  
DT Utility  
FS GRANTED  
LN.CNT 1519  
INCL INCLM: 424/426.000  
INCLS: 424/457.000; 424/462.000; 424/486.000; 424/489.000; 424/490.000;  
424/463.000; 514/772.300; 514/963.000  
NCL NCLM: 424/426.000  
NCLS: 424/457.000; 424/462.000; 424/463.000; 424/486.000; 424/489.000;  
424/490.000; 514/772.300; 514/963.000  
IC [7]  
ICM: A61F002-00  
ICS: A61K009-52; A61K009-58; A61K009-14; A61K047-30  
EXF 424/426; 424/501; 424/457; 424/462; 424/463; 424/486; 424/489; 514/963;  
514/772.3  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L9 ANSWER 3 OF 3 USPATFULL  
AN 2001:188689 USPATFULL  
TI Taxol-like protein (TALP) and process for preparing the same  
IN Hwang, Byung-Doo, Taejon, Korea, Republic of  
Lim, Kyu, Taejon, Korea, Republic of  
Kwak, Sang-Tae, Chungcheongbuk-do, Korea, Republic of

Kweon, Gi-Ryang, Daegu, Korea, Republic of  
Yoon, Wan-Hee, Taejon, Korea, Republic of  
PA Byung-Doo Hwang (non-U.S. corporation)  
PI US 2001034322 A1 20011025  
AI US 2001-799409 A1 20010305 (9)  
RLI Continuation of Ser. No. US 1999-269146, filed on 12 Mar 1999, PENDING  
A 371 of International Ser. No. WO 1996-KR158, filed on 12 Sep 1996,  
UNKNOWN  
DT Utility  
FS APPLICATION  
LN.CNT 665  
INCL INCLM: 514/002.000  
INCLS: 435/184.000  
NCL NCLM: 514/002.000  
NCLS: 435/184.000  
IC [7]  
ICM: C12N009-99  
ICS: A61K038-57  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

=>

L16 ANSWER 1 OF 232 USPATFULL

ACCESSION NUMBER: 2002:209101 USPATFULL  
TITLE: **Hydroxyapatite**-targeting poly (ethylene glycol) and related polymers  
INVENTOR(S): Roberts, Michael James, Madison, AL, United States  
Kozlowski, Antoni, Huntsville, AL, United States  
PATENT ASSIGNEE(S): Shearwater Corporation, Huntsville, AL, United States  
(U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6436386	B1	20020820
APPLICATION INFO.:	US 2000-712536		20001114 (9)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	GRANTED		
PRIMARY EXAMINER:	Page, Thurman K.		
ASSISTANT EXAMINER:	Fubara, Blessing		
LEGAL REPRESENTATIVE:	Alston & Bird LLP		
NUMBER OF CLAIMS:	80		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	0 Drawing Figure(s); 0 Drawing Page(s)		
LINE COUNT:	1412		

OF 232 USPATFULL  
ACCESSION NUMBER:  
TITLE: 2002:157058 USPATFULL  
21784, a novel human calcium channel family member and  
uses thereof  
INVENTOR(S) : Curtis, Rory A.J., Southborough, MA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002081657	A1	20020627
APPLICATION INFO.:	US 2001-875423	A1	20010605 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-209257P	20000605 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	LOUIS MYERS, FISH & RICHARDSON P.C., 225 Franklin Street, Boston, MA, 02110-2804	

NUMBER OF CLAIMS: 31  
EXEMPLARY CLAIM: 1  
NUMBER OF DRAWINGS: 4 Drawing Page(s)  
LINE COUNT: 5663

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides isolated nucleic acids molecules, designated  
21784 nucleic acid molecules, which encode novel calcium channel

L19 ANSWER 47 OF 132 USPATFULL  
ACCESSION NUMBER: 2002:136533 USPATFULL  
TITLE: Method for delivering bioactive agents using  
cochleates  
INVENTOR(S): Unger, Evan C., Tucson, AZ, United States  
PATENT ASSIGNEE(S): Imarx Therapeutics, Inc., Tucson, AZ, United States  
(U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6403056	B1	20020611
APPLICATION INFO.:	US 2000-540448		20000331 (9)
RELATED APPLN. INFO.:	Division of Ser. No. US 1997-925353, filed on 8 Sep 1997, now patented, Pat. No. US 6120751 Continuation-in-part of Ser. No. US 1997-823791, filed on 21 Mar 1997, now patented, Pat. No. US 6143276 Continuation-in-part of Ser. No. US 1997-851780, filed on 6 May 1997, now patented, Pat. No. US 6090800 Continuation-in-part of Ser. No. US 1997-877826, filed on 18 Jun 1997 Continuation-in-part of Ser. No. US 1997-887215, filed on 2 Jul 1997, now patented, Pat. No. US 6028066		

DOCUMENT TYPE: Utility  
FILE SEGMENT: GRANTED  
PRIMARY EXAMINER: Hartley, Michael G.  
LEGAL REPRESENTATIVE: Woodcock Washburn LLP  
NUMBER OF CLAIMS: 63  
EXEMPLARY CLAIM: 1  
NUMBER OF DRAWINGS: 8 Drawing Figure(s); 4 Drawing Page(s)  
LINE COUNT: 6445

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention is directed to charged lipids, compositions comprising charged lipids, and the use of these compositions in drug delivery, targeted drug delivery, therapeutic imaging and diagnostic imaging, as well as their use as contrast agents.

L19 ANSWER 130 OF 132      USPATFULL  
ACCESSION NUMBER:      96:111166 USPATFULL  
TITLE:      Therapeutic drug delivery systems  
INVENTOR(S) :      Unger, Evan C., Tucson, AZ, United States  
                  Fritz, Thomas A., Tucson, AZ, United States  
                  Matsunaga, Terry, Tucson, AZ, United States  
                  Ramaswami, VaradaRajan, Tucson, AZ, United States  
                  Yellowhair, David, Tucson, AZ, United States  
                  Wu, Guanli, Tucson, AZ, United States  
PATENT ASSIGNEE(S) :      ImaRx Pharmaceutical Corp., Tucson, AZ, United States  
(U.S. corporation)

NUMBER	KIND	DATE
-----	-----	-----
US 5580575		19961203
US 1993-76250		19930611 (8)
Continuation-in-part of Ser. No. US 1991-716899, filed on 18 Jun 1991, now abandoned And a continuation-in-part of Ser. No. US 1991-717084, filed on 18 Jun 1991, now patented, Pat. No. US 5228446		

which

is a continuation-in-part of Ser. No. US 1990-569828,  
filed on 20 Aug 1990, now patented, Pat. No. US

5088499

which is a continuation-in-part of Ser. No. US  
1989-455707, filed on 22 Dec 1989, now abandoned

DOCUMENT TYPE:

Utility

FILE SEGMENT:

Granted

PRIMARY EXAMINER:

Kishore, Gollamudi S.

LEGAL REPRESENTATIVE:

Woodcock Washburn Kurtz Mackiewicz & Norris

NUMBER OF CLAIMS:

17

EXEMPLARY CLAIM:

1

NUMBER OF DRAWINGS:      32 Drawing Figure(s); 21 Drawing Page(s)

LINE COUNT:      2932

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB      Therapeutic drug delivery systems comprising

L19 ANSWER 128 OF 132

USPATFULL

ACCESSION NUMBER:

97:122866 USPATFULL

TITLE:

Thermosensitive biodegradable polymers based on  
poly(ether-ester)block copolymers

INVENTOR(S) :

Cha, Younsik, Salt Lake City, UT, United States

Choi, Young Kweon, Salt Lake City, UT, United States

Bae, You Han, Kwangju, Korea, Republic of

PATENT ASSIGNEE(S) :

Macromed, Inc., Salt Lake City, UT, United States

(U.S.

corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5702717		19971230
APPLICATION INFO.:	US 1995-548185		19951025 (8)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		

PRIMARY EXAMINER:

Dean, Ralph H.

LEGAL REPRESENTATIVE:

Thorpe, North & Western, L.L.P.

NUMBER OF CLAIMS:

32

EXEMPLARY CLAIM:

1

LINE COUNT:

1194

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A system and method for the parenteral delivery of a drug in a  
biodegradable polymeric matrix to a warm blooded animal as a liquid

with

the resultant formation of a gel depot for the controlled  
release of the drug. The system comprises an injectable  
biodegradable block copolymeric drug delivery liquid  
having reverse thermal gelation properties. The delivery liquid is an  
aqueous solution having dissolved or dispersed therein an effective  
amount of a drug intimately contained in a biodegradable block

copolymer

matrix. The copolymer has a reverse gelation temperature below the body  
temperature of the animal to which it is administered and is made up of  
(i) a hydrophobic A polymer block comprising a member selected from the  
group consisting of poly(.alpha.-hydroxy acids) and poly(ethylen

L19 ANSWER 99 OF 132 USPATFULL

ACCESSION NUMBER: 2001:188729 USPATFULL

TITLE: WATER SOLUBLE **PACLITAXEL** DERIVATIVES

INVENTOR(S) : LI, CHUN, MISSOURI CITY, TX, United States

WALLACE, SIDNEY, HOUSTON, TX, United States  
YU, DONG-FANG, HOUSTON, TX, United States  
YANG, DAVID J., SUGAR LAND, TX, United States

NUMBER	KIND	DATE
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PATENT INFORMATION:	US 2001034363	A1 20011025
APPLICATION INFO.:	US 1998-50662	A1 19980330 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1997-815104, filed on 11 Mar 1997, GRANTED, Pat. No. US 5977163	

NUMBER	DATE
--------	------

PRIORITY INFORMATION:	US 1996-13184P	19960312 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	RONALD J. KAMIS, FOLEY & LARDNER, 3000 K STREET N.W., SUITE 500, WASHINGTON, DC, 20007-5109	
NUMBER OF CLAIMS:	51	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	17 Drawing Page(s)	
LINE COUNT:	2480	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Disclosed are water soluble compositions of **paclitaxel** and docetaxel formed by conjugating the **paclitaxel** or docetaxel to a water soluble polymer such as poly-glutamic acid, poly-aspartic acid or poly-lysine. Also disclosed are methods of using the compositions for treatment of tumors, auto-immune disorders such as rheumatoid arthritis. Other embodiments include the coating of implantable stents for prevention of restenosis.

132 USPATFULL  
ACCESSION NUMBER: 2002:54341 USPATFULL  
TITLE: Delivery of therapeutic biologicals from implantable tissue matrices  
INVENTOR(S): MacLaughlin, David T., Saugus, MA, UNITED STATES  
Vacanti, Joseph P., Winchester, MA, UNITED STATES  
Donahoe, Patricia K., Boston, MA, UNITED STATES  
Masiakos, Peter T., Boston, MA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002031500	A1	20020314
APPLICATION INFO.:	US 2001-770339	A1	20010126 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-178842P	20000127 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Patrea L. Pabst, Arnall Golden & Gregory, LLP, 2800 One	
	Atlanta Center, 1201 West Peachtree Street, Atlanta, GA, 30309-3450	
NUMBER OF CLAIMS:	23	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	3 Drawing Page(s)	
LINE COUNT:	1457	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Normal cells, such as fibroblasts or other tissue or organ cell types, are genetically engineered to express biologically active, therapeutic agents, such as proteins that are normally produced in small amounts, for example, MIS, or other members of the TGF-beta family

Herceptin.TM.,  
interferons, and anti-angiogenic factors. These cells are seeded into a matrix for implantation into the patient to be treated. Cells may also be engineered to include a lethal gene, so that implanted cells can be destroyed once treatment is completed. Cells can be implanted in a variety of different matrices. In a preferred embodiment, these matrices are implantable and biodegradable over a period of time equal to or less than the expected period of treatment, when cells engraft to form a